4/4/17 OK

CERTIFICATION

SDG No:

1701420R1-1

Laboratory:

Eurofins, Folson, CA

Site:

BMSMC

Matrix:

Air

SUMMARY:

Air samples (Table 1) were collected on the BMSMC facility. The BMSMC facility is located in Humacao, PR. Samples were taken January 22 and 24, 2017 and were analyzed in Eurofins Laboratory of Folson, California that reported the data under SDG No.: 1701420R1-1. The sample results were assessed according to USEPA the documents described in the following order of precedence: QC criteria from "Compendium Method TO-17. Determination of Volatile Organic Compounds (VOCs) In Ambient Air Using Active Sampling Onto Sorbent Tubes (modified), January, 1999"; In addition the following guideline is employed for set up of the GC/MS analytical system including column selection, MS tune requirements, calibration protocols, etc., as per TO-17 method requirements: USEPA Hazardous Waste Support Branch. Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #6. June, 2014). The analyses performed are shown in Table 1. Individual data review worksheets are enclosed for each target analyte group. The data sample summary form shows analytes results that were qualified.

In summary, the results are valid and can be used for decision making purposes.

Table 1. Samples analyzed and analysis performed

SAMPLE ID	SAMPLE DESCRIPTION	MATRIX	ANALYSIS PERFORMED
1701420R1-01A	B18-1IA-012117	Air	Naphthalene
1701420R1-02A	B18-1IAD-012117	Air	Naphthalene
1701420R1-03A	B181IA-2-012117	Air	Naphthalene
1701420R1-04A	B181IA-3-012117	Air	Naphthalene
1701420R1-05A	B181IA-4-012117	Air	Naphthalene
1701420R1-06A	B181 A-5-012117	Air	Naphthalene
1701420R1-07A	B301IA-1-012117	Air	Naphthalene
1701420R1-08A	B301IA-2-012117	Air	Naphthalene
1701420R1-09A	B301IA-3-012117	Air	Naphthalene
1701420R1-10A	B301IA-4-012117	Air	Naphthalene
1701420R1-11A	B301IA-4D-012117	Air	Naphthalene
1701420R1-12A	B301IA-5-012117	Air	Naphthalene
1701420R1-13A	B1830AA-012117	Air	Naphthalene
1701420R1-14A	B1830FB-012217	Air	Naphthalene
1701420R1-15A	B8IA-2-012317	Air	Naphthalene
1701420R1-16A	B8IA-2D-012317	Air	Naphthalene
1701420R1-17A	B8AA-012317	Air	Naphthalene
1701420R1-19A	B8SS-2D-012417	Air	Naphthalene

Reviewer Name:

Rafael Infante

Chemist License 1888

Signature:

Date:

March 27, 2017



Naphthalene-d8

Air Toxics

Client Sample ID: B18-1IA-012117 Lab ID#: 1701420R1-01A

EPA METHOD TO-17

File Name: Dil. Factor: Compound	18013112R1 Date of Extraction: NADate of Collection: 1/22/17 6 1.00 Date of Analysis: 1/31/17 03:			
	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0.060	3.4	0.20
Air Sample Volume(L): 16.8 Container Type: TO-17 VI Tube				
Surrogates		%Recovery		Method Limits

121



50-150



Client Sample ID: B18-11AD-012117 Lab ID#: 1701420R1-02A

Dil. Factor:	1.00	Date	tion: NADate of Collection: 1/22/17 6:09:00 PN Date of Analysis: 1/31/17 04:39 PM		
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.058	3.1	0.18	
Air Sample Volume(L): 17.3 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		107		50-150	





Client Sample ID: B18IA-2-012117 Lab ID#: 1701420R1-03A

File Name: Dil. Factor:	18013114R1 Date of Extraction: NADate of Collec 1.00 Date of Analys			rsis: 1/31/17 0 <u>5:20 PM</u>	
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.065	5.0	0.32	
Air Sample Volume(L): 15.4 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		108		50-150	





Client Sample ID: B18IA-3-012117 Lab ID#: 1701420R1-04A

Dil. Factor:	18013115R1 Date of Extraction: NADate of Collection: 1.00 Date of Analysis: 1				
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.059	1.6	0.096	
Air Sample Volume(L): 17.0 Container Type: TO-17 VI Tube					
••				Method	
Surrogates		%Recovery		Limits	
Nanhthalene-d8	·	106		50-150	





Client Sample ID: B18IA-4-012117 Lab ID#: 1701420R1-05A

File Name: Dil. Factor: Compound	18013116R1 Date of Extraction: NADate of Collection: 1/22/17 6:29:00 PM 1.00 Date of Analysis: 1/31/17 06:41 PM				
	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.066	2.5	0.17	
Air Sample Volume(L): 15.2 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		123		50-150	





Client Sample ID: B18IA-5-012117 Lab ID#: 1701420R1-06A

File Name:	18013117R1 Date of	f Extraction: NADate	of Collection: 1/2	2/17 6:14:00 PM
Dil. Factor:	1.00	of Analysis: 1/31/	17 07:21 PM	
	Rpt. Limit	Rpt. Limit	Amount	Amount
Compound	(ng)	(ug/m3)	(ng)	(ug/m3)
Naphthalene	1.0	0.058	3.0	0.17
Air Sample Volume(L): 17.3				
Container Type: TO-17 VI Tube				
				Method
Surrogates		%Recovery		Limits
Naphthalene-d8		95		50-150





Client Sample ID: B30IA-1-012117 Lab ID#: 1701420R1-07A

Dil. Factor: Compound	18013118R1 Date of Extraction: Nate of Collection: 1.00 Date of Analysis: 1				
	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1,0	0.059	2.0	0.12	
Air Sample Volume(L): 17.0 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		128		50-150	





Client Sample ID: B30IA-2-012117 Lab ID#: 1701420R1-08A

EPA METHOD TO-17

File Name: Dil. Factor:	18013119R1 D 1.00		Date of Collection: 1/2 Date of Analysis: 1/31	
Compound	Rpt. Limi (ng)		Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0,028	0.94 JQ	0.026 JQ

Air Sample Volume(L): 36.0

J = Estimated value.

Q = The internal standard associated with the analyte exceeded acceptance limits.

Container Type: TO-17 VI Tube

Surrogates	%Recovery	Limits
Naphthalene-d8	57	50-150





Client Sample ID: B30IA-3-012117 Lab ID#: 1701420R1-09A

File Name: Dil. Factor:	18013120R1 Date of Extraction: NADate of Collection: 1/22/17 11:18:00 Al 1.00 Date of Analysis: 1/31/17 10:26 PM			
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0.026	3.1	0.084
Air Sample Volume(L): 37.6 Container Type: TO-17 VI Tube				
Surrogates		%Recovery		Method Limits
Naphthalene-d8		113		50-150





Client Sample ID: B30IA-4-012117 Lab ID#: 1701420R1-10A

File Name: Dil. Factor:	18013121R1 Date of Extraction: NADate of Collection: 1/22/17 6:22:00 PM 1.00 Date of Analysis: 1/31/17 11:07 PM				
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.058	2.1	0.12	
Air Sample Volume(L): 17.3 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		125		50-150	





Client Sample ID: B301A-4D-012117 Lab ID#: 1701420R1-11A

File Name: Dil. Factor:	18013122R1 Date of	of Collection: 1/2: of Analysis: 1/31/	1/22/17 7:00:00 PM /31/17 11:47 PM		
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0,064	2.1	0.13	
Air Sample Volume(L): 15.7 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8	_	116		50-150	





Client Sample ID: B30IA-5-012117 Lab ID#: 1701420R1-12A

Dil. Factor:	1.00	Date	of Analysis: 2/1/1	7 12:28 AM
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0.057	2.6	0.15
Air Sample Volume(L): 17.5 Container Type: TO-17 VI Tube				
oontainer Type. 10-17 VI Tube				Method
Surrogates		%Recovery		Limits
Naphthalene-d8		50		50-150





Client Sample ID: B1830AA-012117 Lab ID#: 1701420R1-13A

File Name: Dil. Factor:	18013124R1 Date o 1.00		NADate of Collection: 1/22/17 6:42:00 PM Date of Analysis: 2/1/17 01:08 AM				
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)			
Naphthalene	1.0	0.055	1,4	0.080			
Air Sample Volume(L): 18.2 Container Type: TO-17 VI Tube							
Surrogates		%Recovery		Method Limits			
Naphthalene-d8		116		50-150			





Client Sample ID: B1830FB-012217 Lab ID#: 1701420R1-14A

EPA METHOD TO-17

 File Name:
 18013111R1
 Date of Extraction: NaDate of Collection: 1/22/17 11:35:00 AM

 Dil. Factor:
 1.00
 Date of Analysis: 1/31/17 03:17 PM

 Compound
 Rpt. Limit (ng)
 Rpt. Limit (ug/m3)
 Amount (ng)
 Amount (ug/m3)

 Naphthalene
 1.0
 0.026
 0.72 J
 0.019 J

Air Sample Volume(L): 37.6

J = Estimated value.

Container Type: TO-17 VI Tube

		Method
Surrogates	%Recovery	Limits
Naphthalene-d8	124	50-150





Client Sample ID: B8IA-2-012317 Lab ID#: 1701420R1-15A

Dil. Factor: Compound	1.00	of Analysis: 2/1/1	1/17 01:49 AM		
	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)	
Naphthalene	1.0	0.058	1.7	0.097	
Air Sample Volume(L): 17.3 Container Type: TO-17 VI Tube					
Surrogates		%Recovery		Method Limits	
Naphthalene-d8		125		50-150	





Client Sample ID: B8IA-2D-012317 Lab ID#: 1701420R1-16A

File Name: Dil. Factor:	18013126R1 Date o 1.00	4/17 12:23:00 Pi 7 02:29 AM		
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0.059	1.8	0.10
Air Sample Volume(L): 17.0 Container Type: TO-17 VI Tube				
Surrogates		%Recovery		Method Limits
Naphthalene-d8		126		50-150





Client Sample ID: B8AA-012317 Lab ID#: 1701420R1-17A

Dil. Factor: Compound	18013127R1 Date o 1.00	f Extraction: NADate Date	of Analysis: 2/1/1	
	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)
Naphthalene	1.0	0.054	1.9	0.10
Air Sample Volume(L): 18.5 Container Type: TO-17 VI Tube				
Surrogates		%Recovery		Method Limits
Naphthalene-d8		82	··· ×	50-150





Client Sample ID: B8SS-2D-012417 Lab ID#: 1701420R1-19A

File Name: Dil. Factor:	18013129R1 Date of Extraction: NADate of Collection: 1/24/17 3:29:00 PM 1.00 Date of Analysis: 2/1/17 04:31 AM								
Compound	Rpt. Limit (ng)	Rpt. Limit (ug/m3)	Amount (ng)	Amount (ug/m3)					
Naphthalene	1,0	2,5	Not Detected	Not Detected					
Air Sample Volume(L): 0.400 Container Type: TO-17 VI Tube									
Surrogates		%Recovery		Method Limits					
Naphthalene-d8		109		50-150					



TO-17 SAMPLE COLLECTION



CHAIN-OF-CUSTODY RECORD

Sample Transportation Notice
Relinquishing signature on this document indicates that sample is being shipped in compliance with all applicable local, State, Federal, national, and international laws, regulations and ordinances of any kind. Eurofins assumes no liability with respect to the collection, handling or shipping of these samples. Relinquishing signature also indicates agreement to hold harmless, defend, and indemnify Eurofins against any claim, demand, or action, of any kind, related to the collection, handling, or shipping of samples. D.O.T. Hotline (800) 467-4922.

180 BLUE RAVINE ROAD, SUITE B **FOLSOM, CA 95630** (916) 985-1000 FAX (916) 985-1020

Page 1 of 2

Project Manager Terry Taylor			Project	et Info:				Around	Reporting		T	ī
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Lab I.D. Field Sample I.D. (Location)	Engraved or Stamped Tube #	Date of Collection (mm/dd/yy)	Start Time (hr : min)	Date of Retrieval (mm/dd/yy)	End Time (hr:min)	Pre-1 Flow I	Rate	Post-Tes	st Volume	Indoor Air	Soil Vapor	Other (
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034 BISTA-2-012112	154190	1/22/17	1040	1/22/17	1800	35	-	35	159	Mic	ם כ	
044 BISIA-3-0,2117	13092	1/21/17	1810	1/22/17	1810	36	,	35	17.0	M	שוב	
054 BISTA-4-012117	132020	1/22/17	1100	1/22/17	1829	34		34	15.2	54 (<u> </u>	
06A BISTA-5-012117	152285	1/21/17	१४१५	1/22/17	1814	36		36	17.3	DE C][10
OFA B30[A-1-0,2117	155 296	1/21/17	1817	1/22/17	1817	36		35	17	X	<u> </u>	םנ
07A B30IA-2-012117	149846	1/21/17	1820	1/22/17	1115	35		36	36][回
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Relinquished by: (signature) Date/Time Received by: (signature)				ne								
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TO-17 SAMPLE COLLECTION

CHAIN-OF-CUSTODY RECORD

Sample Transportation Notice
Relinquishing signature on this document indicates that sample is being shipped in compliance with all applicable local, State, Federal, national, and international laws, regulations and ordinances of any kind. Eurofins assumes no liability with respect to the collection, handling or shipping of these samples. Relinquishing signature also indicates agreement to hold harmless, defend, and indemnify Eurofins against any claim, demand, or action, of any kind, related to the collection, handling, or shipping of samples. D.O.T. Hotline (800) 467-4922.

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Page 2 of 2

Project Mar	nager Terry	Taylor				Project	Info:				Around	Reporting	П	Т	7-	7
Collected b	y: (Print and Sign)	Terry Taylo	L.			- P.O. #					ime: ormal	Units:		-	1	إ
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154	B81A-2-01	2317	149175	1/23/17	12	23	1/23/17	122	3 3.	5	37	17.3	প্র]
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			6.5 <u>SMGT</u>													1

EXECUTIVE NARRATIVE

SDG No:

and the

1701420R1-1

Laboratory:

Eurofins, Folson, CA

Analysis: Location: TO-17

Number of Samples:

18

SUMMARY:

Eighteen (18) samples were analyzed for naphthalene in ambient air following Compendium Method TO-17. The sample results were assessed according to USEPA data validation documents in the following order of precedence: QC criteria from "Compendium Method TO-17. Determination of Volatile Organic Compounds (VOCs) In Ambient Air Using Active Sampling Onto Sorbent Tubes (modified), January, 1999"; In addition the following guideline is employed for the evaluation of the set-up of the GC/MS analytical system including column selection, MS tune requirements, calibration protocols, etc., as per TO-17 method requirements: USEPA Hazardous Waste Support Branch. Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #6. June, 2014). The QC criteria and data validation actions listed on the data review worksheets are from the primary guidance document, unless otherwise noted.

Results are valid and can be used for decision making purposes.

Critical issues:

Major:

None None

Minor:

None

Critical findings:

Major findings:

None

1. Results for sample B8SS-2-012417 were not reported due to significant matrix interference which resulted in extremely poor recovery of the internal standard Bromofluorobenzene. As a result, compounds quantified using Bromofluorobenzene could not be evaluated and were therefore not reported.

Minor findings:

- 1. All samples analyzed within the recommended method holding time. Samples received in good conditions and no receiving discrepancies were observed except the cases described in this document. A Temperature Blank was not included with the shipment. Temperature was measured on a representative sample and was not within 4±2 °C. Coolant in the form of blue ice was present. Analysis proceeded; no action taken professional judgment. The % difference in sample flow rate (beginning/end) was within method performance criteria.
- 2. Naphthalene detected in the laboratory blank at a concentration below the reporting limits. No action taken, professional judgment.

Naphthalene was detected in sample 1701420-14A (Blank tube) analyzed with this data package. No action taken, analyte concentration below the reporting limit.

3. The recovery of internal standard Bromofluorobenzene in sample B30IA-2-012117 was outside control limits. Reanalysis of a back-up tube sample confirmed results. Compounds quantified using the affected internal standard are qualified as estimated (J-) values and are flagged with a 'Q' by the laboratory.

COMMENTS:

Results are valid and can be used for decision making purposes.

Reviewers Name:

Rafael Infante

Chemist License 1888

Rafuel Infant

Signature:

Date:

March 27, 2017

NAPHTHALENE DATA SAMPLE SUMMARY

METHOD:

TO-17

NAP	HTI	НΔ	l FI	NF	- T	\cap	17
IVOL			ᄔᄔ	11	_	v	

Sample ID	Date	Results	Units	Dilution Factor	Lab Flag	Validation	Reportable
1701420R1-01A	1/22/2017	3.4	ng	1.0	Q.	12	Yes
1701420R1-02A	1/22/2017	3.1	ng	1.0	-	-	Yes
1701420R1-03A	1/22/2017	5.0	ng	1.0	2		Yes
1701420R1-04A	1/22/2017	1.6	ng	1.0	-6		Yes
1701420R1-05A	1/22/2017	2.5	ng	1.0	-	20	Yes
1701420R1-06A	1/22/2017	3.0	ng	1.0	-	1.60	Yes
1701420R1-07A	1/22/2017	2.0	ng	1.0	-	-	Yes
1701420R1-08A	1/22/2017	0.94	ng	1.0	JQ	J-	Yes
1701420R1-09A	1/22/2017	3.1	ng	1.0	7	-	Yes
1701420R1-10A	1/22/2017	2.1	ng	1.0	-	-	Yes
1701420R1-11A	1/22/2017	2.1	ng	1.0	-	-	Yes
1701420R1-12A	1/22/2017	2.6	ng	1.0		-	Yes
1701420R1-13A	1/22/2017	1.4	ng	1.0	-	-	Yes
1701420R1-14A	1/22/2017	0.72	ng	1.0	J	J	Yes
1701420R1-15A	1/24/2017	1.7	ng	1.0	7	570	Yes
1701420R1-16A	1/24/2017	1.8	ng	1.0	-	7-2	Yes
1701420R1-17A	1/24/2017	1.9	ng	1.0	47	-	Yes
1701420R1-19A	1/24/2017	1.0	ng	1.0	-	U	Yes

Project	Number:_1701420R1	
Date:_	01/22_&_24/2017	_

The following guidelines for evaluating volatile of actions. This document will assist the reviewer in decision and in better serving the needs of the data USEPA the documents described in the following Method TO-17. Determination of Volatile Organic Conto Sorbent Tubes (modified), January, 1999"; I evaluation of the set-up of the GC/MS analytical systealibration protocols, etc., as per TO-17 method revalidating Air Samples. Volatile Organic Analysis of HW-31. Revision #6. June, 2014). The QC criteria worksheets are from the primary guidance document The hardcopied (laboratory name) _EurofinsAir_	LE ORGANIC PACKAGE Irganics were created to delineate required validation in using professional judgment to make more informed a users. The sample results were assessed according to order of precedence: QC criteria from "Compendium ompounds (VOCs) In Ambient Air Using Active Sampling In addition the following guideline is employed for the stem including column selection, MS tune requirements, quirements: USEPA Hazardous Waste Support Branch of Ambient Air in Canisters by Method TO-15, (SOP # a and data validation actions listed on the data review att, unless otherwise noted. Toxics data package received has been ata summarized. The data review for VOCs included:
Lab. Project/SDG No.:1701420R1 No. of Samples:18	
Trip blank No.:B1830FB-012217 Field blank No.: Equipment blank No.: Field duplicate No.:B18-1IA-012117/B18-1IAB8IA-2-012317/B8IA-2D-	AD-012117;_B30IA-4-012117/B30IA-4D-012117;
X Data CompletenessX Holding TimesX GC/MS TuningX Internal Standard PerformanceX BlanksX Surrogate Recoveries	X Laboratory Control SpikesX Field DuplicatesX CalibrationsX Compound IdentificationsX Compound QuantitationX Quantitation Limits
Overall Comments:Naphthalene_by_m_GC/MS	ethod_TO-17_(modified)_detection_by_full_scan
Definition of Qualifiers: J- Estimated results U- Compound not detected R- Rejected data UJ- Estimated nondetect	
Reviewer: Rafael Defaut	<u></u>
Date: 03/27/2017	

DATA COMPLETENESS

MISSING INFORMATION	DATE LAB. CONTACTED	DATE RECEIVED
1		
- 1		
	>	
	1	

All criteria were metX
Criteria were not met
and/or see below

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

SAMPLE ID	DATE SAMPLED	DATE ANALYZED	> 10% difference in sample flow rate (beginning/end)	ACTION
conditions and no document. A Tempon a representative	receiving discreperature Blank wave sample and woroceeded; no ac	epancies were as not included as not within tion taken profe	observed except the with the shipment. Ten 4±2 °C. Coolant in the essional judgment. % of the coolant in the essional judgment.	amples received in good cases described in this inperature was measured be form of blue ice was difference in sample flow

Criteria

Samples should be refrigerated at <4°C in a clean environment during storage and analyzed within 30 days of sample collection (within one week for limonene, carene, *bis*-chloromethyl ether and labile sulfur or nitrogen containing volatiles). Samples taken on tubes containing multiple sorbent beds should be analyzed as soon as possible after sampling unless it is know in advance that storage will not cause significant sample recovery errors.

Receiving temperature: 14°C

Actions

If holding times are exceeded use professional judgment to qualify positive results and non-detects.

Performance Criteria for the Monitoring Pump

Sampling pump errors can normally be presumed to be in the order of 5% (8). If the pump sampling flow rate measured at the end of sample collection varies more than 10% from that measured at the beginning of sample collection, then that sample is invalidated.

All criteria were metX	
Criteria were not met see below	

GC/MS TUNING

The assessment of the tuning results is to determine if the sample instrumentation is within the standard tuning QC limits. The following actions from the TO-15 compendium method are employed.

Gas Chromatograph/Mass Spectrometer (GC/MS) Instrument Performance Check

Action:

NOTES: This requirement does not apply when samples are analyzed by the Selected Ion Monitoring (SIM) technique.

All mass spectrometer instrument conditions must be identical to those used during the sample analysis. Background subtraction actions resulting in spectral distortions for the sole purpose of meeting the method specifications are contrary to the Quality Assurance (QA) objectives, and are therefore unacceptable.

NOTES: No data should be qualified based on BFB or DFTTP failure. Instances of this should be noted in the narrative.

All ion abundance ratios must be normalized to m/z 95, the nominal base peak, even though the ion abundance of m/z 174 may be up to 120% that of m/z 95.

- 1. If samples are analyzed without a preceding valid instrument performance check, qualify all data in those samples as unusable (R).
- 2. If the laboratory has made minor transcription errors which do not significantly affect the data, the data reviewer should make the necessary corrections on a copy of the form.
- 3. If the laboratory has failed to provide the correct forms or has made significant transcription or calculation errors, the Region's designated representative should contact the laboratory and request corrected data. If the information is not available, the reviewer must use professional judgment to assess the data and notify the Project Officer (PO).
- 4. If ion abundance criteria are not met, professional judgment may be applied to determine to what extent the data may be utilized. When applying professional judgment to this topic, the most important factors to consider are the empirical results that are relatively insensitive to location on the chromatographic profile and the type of instrumentation. Therefore, the critical ion abundance criteria for BFB are the m/z 95/96, 174/175, 174/176, and 176/177 ratios. The relative abundances of m/z 50 and 75 are of lower importance. This issue is more critical for Tentatively Identified Compounds (TICs) than for target analytes.
- 5. Note, in the Data Review Narrative, decisions to use analytical data associated with BFB instrument performance check failures (not meeting contract requirements).
- 6. If the reviewer has reason to believe that instrument performance check criteria were achieved using techniques other than those described in the Compendium method TO-15 entitled "Determination Of Volatile Organic Compounds(VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry(GC/MS)", section 10.4, obtain additional information on the instrument performance checks. If the techniques employed are found to be at variance with the contract requirements, the performance and procedures of the laboratory may merit evaluation.
- 7. Use professional judgment to determine whether associated data should be qualified based on the spectrum of the mass calibration compound.

-			
List	the	samples	affected:
If no, use profes qualified or reject	, ,	ine whether the associated data	should be accepted,
XBFB tunii	ng was performed for every	24 hours of sample analysis.	
XThe BFB	performance results were r	reviewed and found to be within th	e specified criteria.

All criteria were metX
Criteria were not met
and/or see below

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data. The calibration criteria and appropriate actions from the compendium method TO-15 are employed.

Date of initial calibration:_	01/09-10/17
Dates of continuing calibra	ation:01/31/17
Instrument ID numbers:	MSD-18
Matrix/Level:	Air/low

DATE	LAB ID#	FILE	CRITERIA OUT RFs, %RSD, %D, r	COMPOUND	SAMPLES AFFECTED
			rations meet method sp requirements.	ecific requirements. Initial	calibration retention
				,	

The following criteria apply:

Table 5. Initial Calibration Actions for TO-15 Analyses

	Act	on
Criteria for TO-15 Analysis	Detected Associated Compounds	Non-Detected Associated Compounds
RRF < 0.010 (poor response volatile target	J (based on mass	
compounds, Table 4)	spectral	R
RRF < 0.050 (all other volatile target compounds)	identification)	
RRF > 0.010 (poor response volatile target compounds, Table 4) RRF > 0.050 (all other volatile target compounds)	No qualification	
% RSD > 40.0 or < -40.0 (poor response volatile target compounds, Table 4) % RSD > 30.0 or < -30.0 (all other Volatile target compounds)	No qualification	
% RSD < 40.0 and > -40.0 (poor response volatile target compounds. Table 4) % RSD < 30.0 and > -30.0 (all other volatile target compounds)	J	Use professional judgment

Table 6. Continuing Calibration Verification (CCV) Actions for TO-15 Analyses

	Action	
Criteria for CCV	Detected Associated Compounds	Non-Detected Associated Compounds
RRF < 0.010 (poor response volatile target compounds, Table 4) RRF < 0.050 (all other volatile target compounds)	J (based on mass spectral R identification)	
RRF > 0.010 (poor response volatile target compounds, Table 4) RRF > 0.050 (all other volatile target compounds)	No qualification	
%D > 40.0 or < -40.0 (poor response volatile target compounds, Table 4) %D > 30.0 or < -30.0 (all other Volatile target compounds)	j UJ	
%ID < 40.0 and > -40.0 (poor response volatile target compounds, Table 4) %ID < 30.0 and > -30.0 (all other volatile target compounds)	No qualification	

If the % D for daily calibration exceeds -90, use professional judgment to see if non-detects nee to be qualified as unusable "R"

A separate worksheet should be filled for each initial curve

Table 4. TO 15 Volatile Compounds List*

Compound	CAS	Synonyms	
	Number		
Acetone	67-64-1	Dimethyl ketone; Dimethylformaldehyde; 2-Propanone	
Allyl chloride	107-05-1	3-Chloropropene; 3-Chloroprene	
Benzene	71-43-2	Benzol; Benzine	
Benzyl chloride	100-44-7	Chloromethylbenzene; alpha-Chlorotoluene	
Bromodichloromethane	75-27-4	Monobromodichloromethane: Methane-bromodichloro	
Bromoethene	593-60-2	Vinyl bromide; Monobromoethene	
Bromoform	75-25-2	Tribromoethane	
Bromomethane	74-83-9	Methyl bromide; Monobromomethane	
1.3-Butadiene	106-99-0	Biethylene; Erythrene; Pyrrolyene	
Carbon disulfide	75-15-0	Carbon bisulfide; Carbon sulfide	
Carbon tetrachloride	56-23-5	Carbon tet; Tetrachloromethane	
Chlorobenzene	108-90-7	Monochlorobenzene; Chlorobenzol; Benzene chloride	
Chloroethane	75-00-3	Ethyl chloride; Chlorene; Chloryl	
Chloroethene	75-01-4	Vinyl chloride; Ethylene monochloride	
Chloroform	67-66-3	Trichloromethane; Methyltrichloride; Methane trichloride	
Chloromethane	74-87-3	R40; Methyl chloride; Monochloromethane	
Cyclohexane	110-82-7	Hexamethylene; Hexahydrobenzene; Hexanaphthene	
Dibromochloromethane	124-48-1	Chlorodibromomethane	
1,2-Dibromoethane	106-93-4	EDB; Ethylene dibromide	
1.2-Dichtorobenzene	95-50-1	ODB; Chloroben	
1.3-Dichlorobenzene	541-73-l	meta-Dichlorobenzene; m-Phenylenedichloride	
1.4-Dichlorobenzene	106-46-7	para-Dichlorobenzene; Parazene; Santochlor	
1.1-Dichloroethane	75-34-3	Ethylidene chloride; Ethylidene dichloride	
1.2-Dichloroethane	107-06-2	Ethylene dichloride; Glycol dichloride; 1.2-DCA	
1.1-Dichloroethene	75-35-4	1.1-DCE: Vinylidene chloride	
cis-1.2-Dichloroethylene	156-59-2	cis-1.2-DCE; cis-Acetylene dichloride	
trans-1.2-Dichloroethylene	156-60-5	trans-1,2-DCE; trans-Acetylene dichloride	
1,2-Dichtoropropane	78-87-5	Propylene dichloride; Propylene chloride	
cis-1.3-Dichloropropene	10061-01-5	1-Propene,1,3-dichloro-,(z)-; cis-1,3-Dichloro-1-Propene	
trans-1.3-Dichloropropene	10061-02-6	trans-1,3-Dichloro-1-Propene; trans-1,3-Dichloropropylene	
1,4-Dioxane	123-91-1	Diethylene dioxide; Diethylene ether	
Ethyl acetate	141-78-6	Acetic acid ethyl ester: Acetic ether	
Ethylbenzene	100-41-4	Ethylbenzol; Phenylethane	
4-Ethyltoluene	622-96-8	1-Ethyl-4-methyl benzene; p-Methylethylbenzene	
Freon 11 (CCl3F)	75-69-4	Trichlorofluoromethane; Fluorotrichloromethane;	
		Fluorocarbon 11	

Freon 12 (CCI2F2)	75-71-8	Dichlorodifluoromethane; Fluorocarbon 12	
Freon 113 (C2Cl3F3)	76-13-1	1.1.2-Trichloro-1,2.2-trifluoroethane; Fluorocarbon 113; 1.1.2-	
		Trichlorotrifluoroethane	
Freon 114 (C2Cl2F4)	76-14-2	1,2-Dichlorotetrafluoroethane; Halocarbon 114; 1,2-Dichloro-	
		1.1.2.2-tetrafluoroethane	
Heptane	142-82-5	Dipropylmethane; Heptyl hydride	
Hexachlorobutadiene	87-68-3	1.3-Hexachtorobutadiene; Perchlorobutadiene	
Hexane	110-54-3	n-Hexane; Hexyl hydride	
2-Hexanone	591-78-6	Methyl butyl ketone; Butyl methyl ketone; Hexan-2-one	
Isopropyl alcohol	67-63-0	2-Propanol; Isopropanol	
Methylene chloride	75-09-2	Dichloromethane; Methylene dichloride	
Methyl ethyl ketone	78-93-3	MEK; 2-Butanone; Ethyl methyl ketone	
Methyl isobutyl ketone	108-10-1	MIBK; 2-Pentanone; Hexone; Isopropylacetone	
Methyl tert-butyl ether	1634-04-4	MTBE; 2-Methoxy-2-methylpropane; tert-Butyl methyl ether	
Propy lene	115-07-1	Propene; Methylethylene	
Styrene	100-42-5	Vinylbenzene; Phenylethylene	
1.1.2.2-Tetrachloroethane	79-34-5	Tetrachloroethane; Acetylene tetrachloride; Bonoform	
Tetrachloroethene	127-18-4	PCE; PERC; Perchloroethylene; Ethylene tetrachloride; Carbon	
		bichloride; Carbon dichloride	
Tetrahydrofuran	109-99-9	Diethylene oxide; Butylene oxide	
Toluene	108-88-3	Toluol; Methylbenzene	
1.2.4-Trichlorobenzene	120-82-1	1,2,4-Trichlorobenzol	
1.1.1-Trichloroethane	71-55-6	Methyl chloroform; Trichloroethane	
1.1.2-Trichloroethane	79-00-5	beta-Trichloroethane; Ethane trichloride; Vinyl trichloride	
Trichloroethene	79-01-6	TCE; Acetylene trichloride; Ethinyl trichloride	
1.2.4-Trimethylbenzene	95-63-6	Pseudocument Pseudocumol	
1.3.5-Trimethylbenzene	108-67-8	Mesitylene; Trimethylbenzol	
2.2.4-Trimethylpentane	540-84-1	Iso-octane; Isobutyltrimethylmethane	
Vinyl acetate	108-05-4	Acetic acid ethenyl ether; Ethenyl acetate	
p-Xylene	106-42-3	p-Methyltoluene; 1.4-dimethylbenzene	
m-Xylene	108-38-3	m-Methyltoluene; 1.3-dimethylbenzene	
o-Xylene	95-47-6	o-Methyltoluene; 1.2-Dimethylbenzene	

^{*}Laboratories use different sets and subsets of analytes on as needed basis.

NOTES:

Compounds in bold italicized letters may have poor GCMS response. These poor response compounds are evaluated using more relaxed relative response factor criteria as stated below.

Note: Naphthalene does not have poor GCMS response. Calibration criteria: RRF > 0.05 and % difference in the continuing calibration verification < 30 %.

All criteria were met _X
Criteria were not met
and/or see below

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory	þ	lan	ıks
------------	---	-----	-----

DATE ANALYZED	LAB ID	LEVEL/ MATRIX	COMPOUND	CONCENTRATION UNITS
			criteria_except_for_the Naphthalene	•
		··· <u>-</u>		

Note: Naphthalene detected in the laboratory blank at a concentration below the reporting limits. No action taken, professional judgment.

Field blanks

Field blanks are the same as laboratory blanks except that they are transported to and from the monitoring site, are uncapped and immediately resealed at the monitoring site, but do not actually have air pumped through them. One field blank tube is taken for every ten sampled tubes on a monitoring exercise.

Criteria:

If the same profile/pattern of VOCs is observed on the field blanks as on the sampled tubes and if the level of these components is 5% or more of the sampled volatiles, careful attention must be paid to the method of sealing the tubes and other storage procedures in future studies. If the profile of volatiles on the field blanks matches that of the sampled tubes and if the areas of the peaks on the field blank are 10% or more of sampled tube levels, the sampled tube data are invalidated.

DATE ANALYZED	LAB ID	LEVEL/ MATRIX	COMPOUND	CONCENTRATION UNITS
			1420R1-14A_(Blank_tu tration_below_the_repo	be)_analyzed_with_this_data orting_limit

Note:

All criteria were metX
Criteria were not met
and/or see below

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately

Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

CONTAMINATION SOURCE/LEVEL	COMPOUND	CONC/UNITS	AL/UNITS	SQL	AFFECTED SAMPLES
				-	
The second second					
To the same of the					

All criteria were met _	X_
Criteria were not met	
and/or see below	_

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery.

Matrix: solid/aqueous

SΔ		_		חו
~ n	na.		_	

SURROGATE COMPOUND

ACTION

1,2- Naphthalene-d8 4-BFB DICHLOROETHANE-d4

Surrogate_recoveries_within_laboratory_control_limits				
		-		
QC Limits* (Air)				
LL_to_ULto	toto			

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

QUALITY	%R < 10%	%R = 10% - LL	%R > UL
Positive results	J	J	J
Nondetects results	R	UJ	Accept

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

All criteria were metX
Criteria were not met
and/or see below

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

1. LCS Recoveries Criteria

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

	LCS ID	COMPOUND	% R	QC LIMIT
		e)_analyzed_in_this_data_p	oackage;_%_recoverie	es_and_RPD
within_la	boratory_contro	l_limits		

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit
 - If QC limits are not available, use limits of 70 130 %.

Actions:

Table 9. LCS/LCSD Actions for TO-15 Analyses

	Action		
Criteria	Detected Associated Compounds	Non-detected Associated Compounds	
Percent recovery Criteria			
%R > Upper Acceptance Limit (>130%)	J	No qualification	
%R in the acceptable range, 70-130%	No qu	alification	
%R < Lower Acceptance Limit (< 70 %)	J	UJ	
%R < 50%	J	R	
Lower Acceptance Limit \leq %R \leq Upper Acceptance Limit	No qualification		
Relative Percent Difference Criteria		<u> </u>	
% RPD ≤ 25%	No qualification		
% RPD > 25 %	J	ŲJ	

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

I

			All criteria were Criteria were and/or see be	
X.	LABORATORY	//FIELD DUPLICATE PRECISION		
	Sample IDs: Sample IDs: Sample IDs: Sample IDs:	_ B18-1IA-012117/B18-1IAD-012117_(field) _ B30IA-4-012117/B30IA-4D-012117_(field) _ B8IA-2-012317/B8IA-2D-012317_(field) _ LCS/LCSD_(laboratory)	Matrix: Matrix: Matrix: Matrix:	

Field/laboratory duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information.

Suggested criteria: RPD \pm 50% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

COMPOUND	SQL	SAMPLE CONC.	DUPLICATE CONC.	RPD	ACTION
	-				

Note: Laboratory/field duplicates analyzed as part of this data set. Laboratory duplicate were within method performance criteria.

Field duplicates RPD are within method performance criteria.

Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

All criteria were met	_
Criteria were not met	
and/or see belowX_	_

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within ± 20 seconds of the IS area in the associated calibration standard.

DATE	SAMPLE ID	IS OUT	IS AREA	ACCEPTABLE RANGE	ACTION
	andard_area_and_reation_standards_exc				
_01/31/17	_B30IA-2-012117	BFB	39718241 		_Qualify_result n_sample_(J-)
					

Table 10. Internal Standard Actions for TO-15 Analyses

	Action		
Criteria	Detected Associated Compounds*	Non-detected Associated Compounds*	
Area counts > 140% of CCV or mid-point standard from initial calibration)	J-	No qualification	
Area counts < 60% of CCV or mid-point standard from initial calibration)	J+	R	
Area counts ≥ 60% but ≤ 140% of CCV or mid-point standard from initial calibration)	No qualification		
RT difference > 20.0 seconds between samples CCV or mid- point standard from initial calibration)	R*		
RT difference < 20.0 seconds between samples and CCV or mid-point standard from initial calibration)	No qualification		

Examine the chromatographic profile for that sample to determine if any false positives or negatives exist. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for that sample fraction. Detects should not need to be qualified as unusable (R) if the mass spectral criteria are met.

Note:

Actions:

The recovery of internal standard Bromofluorobenzene in sample B30IA-2-012117 was outside control limits. Reanalysis of a back-up tube sample confirmed results. Compounds quantified using the affected internal standard are qualified as estimated values (J-) and are flagged with a 'Q' by the laboratory.

Results for sample 1701420-18A (B8SS-2-012417) were not reported due to significant matrix interference which resulted in extremely poor recovery of the internal standard Bromofluorobenzene. As a result, compounds quantified using Bromofluorobenzene could not be evaluated and were therefore not reported.

All criteria were met _X	
Criteria were not met	
and/or see below	

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

1701420R1-01A

Naphthalene

RF = 1.51236

 $[\]=(280008)(36)/(1952676)(1.51236)$

= 3.413 ng OK

All criteria were metX
Criteria were not met
and/or see below

XII. QUANTITATION LIMITS

A. Dilution performed

SAMPLE ID	DILUTION FACTOR	REASONS FOR DILUTION
No dilution perf	formed.	
		F-10 (
	<u> </u>	
		The state of the s
	7	

System Performance

Action:

Use professional judgment to qualify the data if it is determined that system performance has degraded during sample analyses. Note, for Laboratory Project Officer (PO) action, any degradation of system performance which significantly affected the data.

Note:

Overall Assessment of Data

Action:

1. Use professional judgment to determine if there is any need to qualify data which were not qualified based on the Quality Control (QC) criteria previously discussed.

2. Write a brief narrative to give the user an indication of the analytical limitations of the data. Note, for Laboratory Project Officer (PO) action, any inconsistency of the data with the Sample Delivery Group (SDG) Narrative. If sufficient information on the intended use and required quality of the data is available, the reviewer should include their assessment of the usability of the data within the given context. This may be used as part of a formal Data Quality Assessment (DQA).

Overall assessment of the data:

Results are valid; the data can be used for

decision making purposes.